

Substitute
Spec

91787342
JC02 Rec'd PCT/PTO 16 MAR 2001

INFUSION PUMP COMPRISING A COMPUTER FOR CALCULATING
THE RESPECTIVE MAXIMUM PERMISSIBLE DOSAGE

Prior Applications

This application is a §371 U.S. National Phase application
which bases priority on International Application No.
PCT/DE99/03000, filed September 20, 1999, which in turn bases
priority on German Application No. DE 198 42 722.0 filed
September 18, 1998.

10

Background of the Invention

1. Field of Invention

The invention relates to an infusion pump for the delivery
of a quantity of medicament to the body of a patient
determinable by means of an electronic control device, the pump
being provided with a computer for calculating the maximum
permitted delivery quantity as a function of the previously
delivered quantity and a blocking device for preventing further
medicament delivery on exceeding a predetermined, permitted
maximum value.

20

2. Description of the Prior Art

Such infusion pumps are used for supplying a patient with
a medicament over a long time period and the medicament
quantity continuously delivered by the infusion pump
corresponding to the needs of the patient can be adjusted.

25

DE 33 90 462 C2 discloses an implantable infusion pump

Please enter
this specification
Nagay
after

FILED: 2000-03-16

equipped with a computer, which determines the medicament quantity delivered over a "sliding time window/slot," e.g. over three hours and blocks further delivery if the quantity delivered over this time period exceeds a maximum value.

5 However, this procedure is inadequate and the sliding time window length random. In many cases, such as e.g. with an attack of pain, it is necessary to briefly considerably raise the quantity of active substance to be delivered by the pump in order to rapidly raise the active substance level. However, 10 whereas a quantity distributed over three hours can be tolerated, this can prove toxic when administered over three minutes. However, an infusion rate allowed when distributed over three hours, can prove toxic or even lethal when the administration extends beyond three hours. This problem cannot 15 be solved with the "sliding time window."

 The problem of the invention is to provide an implantable infusion pump making it possible to reliably determine in each case the allowed delivery quantity.

Summary of the Invention

20 According to the invention this problem is solved in that the computer determines the quantity or concentration of the active substance in the body of the patient on the basis of the medicament quantity delivered and its breaking down in the body

and compares it with the predetermined maximum value.

A preferred embodiment is characterized in that the computer is provided with a memory storing a quantity resulting from the adding up of the delivered quantity in each case and a subtraction of the percentage of the quantity entered in the memory resulting from the expected breaking down of the medicament in the body, as well as a comparator which constantly compares the quantity entered in the memory with the predetermined, permitted maximum value.

The maximum value at which blocking takes place is consequently not, as in the case of the prior art, a value averaged out over a given time window, but is in the form of the integral reduced by the amount resulting from the half-life of the medicament over the total quantity delivered.

The computer is preferably provided with a device which, either with a time interval predetermined in accordance with the expected breaking down of the medicament in the body, brings about the subtraction of a specific percentage of the quantity entered in the memory, or in the case of a fixed, predetermined time intervals brings about the subtraction of a percentage of the quantity entered in the memory corresponding to the expected breaking down of the medicament.

In the case of the proposed construction of the infusion

5 pump it is ensured that the administration of the medicament,
which is brought about by means of the control device by the
doctor or optionally also the patient, does not exceed a
maximum permitted value.

5 In order to adjust the device for a given patient, it is
merely necessary to input the half-life of the medicament to be
administered and the individually permitted maximum value
(toxic threshold).

10 It is obvious that the device must also be programmed in
such a way that the lower minimum value (action threshold) is
maintained.

15 The pump can be an implantable infusion pump. It is also
possible to place the computer (or an additional, parallel-
operating computer) in an external control device. It is
possible for a bolus administration (namely an infusion of the
medicament which in the case of long-term administration would
lead to the toxic threshold being exceeded) only being possible
in the case of electromagnetic coupling with the control
device.

20 Description of the Drawings

The invention is described in greater detail hereinafter
relative to the drawing wherein:

Fig. 1 shows, in the lower graph, an infusion profile and,

in the upper graph, the pattern resulting from this infusion profile of the quantity entered in the memory, the expected quantity (and therefore the concentration) of the active substance in the body of the patient, as well as the predetermined, permitted maximum value (threshold S).

Detailed Description of the Preferred Embodiment

In the case of the infusion profile shown there is initially a long-term administration with a relatively low infusion rate. As from time t_1 to t_2 (caused by the patient or doctor) a first bolus administration takes place, i.e. a brief administration with a high infusion rate, such as is e.g. necessary if the patient suffers an acute attack. At time t_3 switching to a higher infusion rate takes place. At time t_4 , using the control device, the administration of a bolus is brought about which, on reaching the predetermined threshold S is prematurely stopped at time t_5 by the computer. At time t_6 the user attempts to set a bolus administration, which is stopped at time t_6 because the threshold S has been reached.

The path of the active substance concentration in the body of the patient resulting from this infusion profile and which is essentially proportional to the active substance quantity present in the body is shown in the lower graph.

The pattern of the active substance concentration is

represented by a time integral over the infused quantity,
reduced by the breaking down resulting from the half-life of
the substance, i.e. as a function with a linear term determined
by medicament administration and a negative exponential term
determined by the medicament braking down rate.

In the drawing, this leads up to time t_1 to a constant
path, because here the quantity supplied precisely corresponds
to the quantity broken down by the body. The administration of
the bolus at time t_1 leads to a steep rise in the active
substance concentration. At the end of bolus administration at
time t_2 the concentration continuously drops, because the
supplied active substance quantity is lower than the broken
down quantity. After doubling the infusion rate at time t_3 the
concentration constantly rises, but with a shallower rise.

The bringing about of a further bolus administration
through the user or doctor at time t_4 leads to a concentration
rise up to the threshold at time t_5 , which at time t_6 leads to
an automatic termination of bolus administration by the
computer. The attempt at time t_7 to bring about a further
bolus administration is immediately prevented by the computer
due to the immediate reaching of the threshold.

The path of the active substance concentration is
simulated in the computer of the implantable infusion pump

(which can also be located in the control device).

In predetermined time intervals, e.g. every 10 sec, the quantity entered in the memory of the infusion pump is increased by a quantity corresponding to the amount delivered by the infusion pump in this time period. Furthermore, a mathematically determined percentage of the quantity entered in the memory is subtracted from the half-life of the delivered medicament, the resulting quantity is stored as the actual value. Alternatively, in time intervals given by the half-life (i.e. more frequently with a shorter half-life and less frequently with a longer half-life), the amount delivered in this time period can be summed and a fixed quantity subtracted.

The value entered in the memory consequently always corresponds (due to the not precisely determinable half-life this is naturally only approximately) to the actual amount in each case or concentration of the active substance in the body of the patient, whilst taking account of the breaking down thereof.